

Section C

510(k) Summary

"This summary of 510(k) safety and effectiveness information is being submitted in accordance with the requirements of SMDA 1990 and 21 CFR 807.92."

"The assigned 510(k) number is: K130800" (applicant leave blank)

Premarket Notification [510(k)] Summary

[(a)(1)]. The summary contains on the first page, preferably on your letterhead paper, the submitter's name, address, phone and fax numbers, name of contact person, and date the summary was prepared :

Submitter's name : Kaixuan Plastic Co., Ltd.
 Submitter's address : No.44, Zhonghua St. Shijiazhuang, 050000, China
 Phone number : (86) 311-67699828
 Fax number : (86) 311-67699906
 Name of contact person: Mr. Zhou Rui
 Date the summary was prepared: 2014-02-07

[(a)(2)]. The name of the device, including the trade or proprietary name if applicable, the common or usual name, and the classification name, if known

Device Name: Powder Free Vinyl Patient Examination Gloves, Clear (non-colored)
 Proprietary/Trade name: Powder Free Vinyl Patient Examination Gloves, Clear (non-colored)
 Common Name: Patient examination glove
 Classification Name: Patient examination glove
 Device Classification: I
 Regulation Number: 21 CFR 880.6250
 Panel: General Hospital (80)
 Product Code: LYZ

[(a)(3)]. An identification of the legally marketed device to which your firm is claiming substantial equivalence .

Class I* Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) that meets all of the requirements of ASTM standard D 5250-06 (Reaffirmation 2011).

Predicate device : Powder free Vinyl Patient Examination Gloves, Clear(Non-colored), Shijiazhuang Fuguan Plastic Products Co., Ltd. K032908 .

[(a)(4)] A description of the device

Device Description : Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) that meets all of the requirements of ASTM standard D 5250-06(Reaffirmation 2011).

-- How the device functions:

PVC films form a barrier to body fluids and bloodborne Pathogens

-- Scientific concepts that form the basis for the device

The PVC rubber is water tight under normal conditions of use. Its tensile properties cause it to conform to the hand, allowing movements necessary for a medical procedure.

-- Physical and performance characteristics such as design, materials and physical properties:

PVC gloves are known to create a barrier to bloodborne pathogens and body fluids. ASTM conforming tensile properties create a glove that is strong and flexible. The glove is manufactured in accordance with the requirements of ASTM D5250 and ASTM D5151 requirements.

[(a)(5)] The summary describes the intended use of the device

Device Intended Use: Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.

[(a)(6)] A summary of the technological characteristics of new device compared to the predicate device.

The Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) non sterile are summarized with the following technological characteristics compared to ASTM or equivalent standard.

Features & Description	Predicate Device	Medical Glove Guidance Manual(1661)	Subject Device	Result of Comparison
Company	Shijiazhuang Fuguan Plastic Products Co., Ltd.		Kaixuan Plastic Co., Ltd.	--
510(K) Number	K032908		K130800	
Product name	Powder Free Vinyl Patient Examination Gloves, Clear (Non-colored)		Powder Free Vinyl Patient Examination Gloves, Clear (non-colored)	Same
Product Code	LYZ	LYZ	LYZ	Same
Size	Small/ Medium/ Large/X large		Small/ Medium/ Large/X large	Substantially equivalent
Intend for use	Powder free Vinyl Patient Examination Gloves, Clear(Non-colored)is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	Powder Free Examination Gloves: A powder-free patient examination glove is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	Powder free Vinyl Patient Examination Gloves, Clear (Non-colored) is a disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.	Substantially equivalent
Device Description and Specifications	Meets ASTM D5250 -06 (Reapproved 2011)	If vinyl: Do the vinyl examination gloves meet all the current specifications listed under ASTM Specification D5250	Meets ASTM D5250 -06 (Reapproved 2011)	Substantially equivalent

		or an equivalent consensus standard?		
Dimensions -- Length	Meets ASTM D5250-06 (Reapproved 2011) ≥230mm min.	ASTM D5250	230mm min for all sizes	Substantially equivalent
Dimensions -- Width	Meets ASTM D5250-06 (Reapproved 2011) Small 80-90 mmr Medium 90-100mm Large 100-110mm X large 110-120 mm		Small 80-85 mm Medium 95-97 mm Large 102-108mm X large 114-118 mm	Substantially equivalent
Dimensions -- Thickness	Meets ASTM D5250-06 (Reapproved 2011) Finger 0.05mm min. Palm 0.08mm min.		Finger 0.05mm min. Palm 0.08mm min.	
Physical Properties	Meets ASTM D5250-06 (Reapproved 2011) Before aging/after aging Elongation ≥300% Tensile Strength≥14MPa	ASTM D5250	Before aging/after aging Elongation ≥300% Tensile Strength≥ 14MPa	Substantially equivalent
Freedom from Pinholes	Meets <ul style="list-style-type: none"> • 21 CFR 800.20 • ASTM D5250-06 (Reapproved 2011) • ASTM D 5151-06 (Reapproved 2011) 	21 CFR 800.20 ASTM D5250 ASTM D 5151	Meets ASTM D5151-06(Reapproved 2011) Holes Inspection Level I AQL2.5	Substantially equivalent
Residual Powder	Meets ASTM D 6124-06(Reapproved 2011)	ASTM D 6124	Meets ASTM D 6124-06(Reapproved 2011) Results generated values below 2mg of residual powder	Substantially equivalent
Compare all materials used to fabricate the devices	PVC	If the glove is made of a polymer or other type of material. identify the material.	PVC	Substantially equivalent
Dusting or Donning Powder:	PU	If a donning lubricant is used, state the composition and include biocompatibility data for the lubricant in an identified attachment; also state the name, manufacturer, and address below	PU	Substantially equivalent
Dusting or Donning Powder: name	PU	Lubricant Generic Name/ Lubricant Brand Name	Surface Coating Agent	Substantially equivalent
Compare performance data supporting substantial equivalence	Meets ASTM D5151-06 (Reapproved 2011) ASTM D5250-06 (Reapproved 2011) ASTM D6124-06 (Reapproved 2011)	At this time FDA recognizes the following standards: Patient Examination Gloves(PVC) ASTM D5151(Detection of Holes in Medical Gloves) ASTM	Meets ASTM D5151-06 (Reapproved 2011) ASTM D5250-06 (Reapproved 2011) ASTM D6124-06 (Reapproved 2011)	Substantially equivalent

		D6124(Residual Powder on Medical Gloves) ASTM D5250(Poly(vinyl chloride) Gloves)		
Single Patient Use	Single Patient Use	Single Patient Use	Single Patient Use	Substantially equivalent
Biocompatibility	SKIN IRRITATION DERMAL and SENSITIZATION STUDIES Meets ISO 10993-10 :2002/Amd.1: 2006	SKIN IRRITATION DERMAL and SENSITIZATION STUDIES ISO 10993-10	The test article was a non-irritant or non-sensitizer SKIN IRRITATION DERMAL and SENSITIZATION STUDIES Meets ISO ISO 10993-10 Third Edition 2010-08-01	Very similar only ISO 10993-10 version updated Substantially equivalent
Labeling for the legally marketed device to which substantial equivalence is claimed.	-Powder Free -Patient Examination Glove -glove color, clear (non-colored) - non sterile -Single Use Only - Manufactured For: - Lot	Chapter 4	-Powder Free -Patient Examination Glove -glove color, clear (non-colored) - non sterile -Single Use Only - Manufactured For: - Lot	Substantially equivalent

[(b)(1)] A brief discussion of the non-clinical submitted, reference, or relied on in the premarket notification submission for a determination of substantial equivalence .

Powder Free Vinyl Patient Examination Gloves, Clear (non-colored)meet requirements per ASTM D5250-06(Reaffirmation 2011), per ASTM D6124-06(Reaffirmation 2011), per 21 CFR 800.20 and ISO 10993-10 Third Edition 2010-08-01.

[(b)(2)] A brief discussion of the clinical submitted, reference, or relied on in the premarket notification submission for a determination of substantial equivalence .

Clinical data is not needed for gloves or for most devices cleared by the 510(k) process.

[(b)(3)] The conclusions drawn from the non-clinical and clinical tests that demonstrate that the device is as safe,as effective, and performed as well or better than the legally marketed device identified in (a)(3).

It can be concluded that the Powder Free Vinyl Patient Examination Gloves meet the ASTM standard or equivalent standard and FDA requirements for waterleak test on pinhole AQL., meet labeling claims and the Powder Free Vinyl Patient Examination Gloves, Clear (non-colored) is as safe, as effective, and performs as well as the predicate device, Powder free Vinyl Patient Examination Gloves, Clear(Non-colored), Shijiazhuang Fuguan Plastic Products Co., Ltd. K032908 .



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
10903 New Hampshire Avenue
Document Control Center – WO66-G609
Silver Spring, MD 20993-0002

February 18, 2014

Kaixuan Plastic Company, Limited
C/O Mr. Chu Xiaolan
Official Correspondent
Room 1606 Bldg 1 Jianxiang Yuan #209
Bei Si Huan Zhong Road, Haidian District
Beijing 100083
CHINA

Re: K130800

Trade/Device Name: Powder Free Vinyl Patient Examination Gloves, Clear (non-colored)
Regulation Number: 21 CFR 880.6250
Regulation Name: Patient Examination Glove
Regulatory Class: I
Product Code: LYZ
Dated: January 2, 2014
Received: January 8, 2014

Dear Mr. Xiaolan:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA).

You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address <http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address <http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

 Tejas Purohit-Sheth, M.D.
Clinical Deputy Director
DAGRID

FOR

Erin I. Keith, M.S.
Acting Director
Division of Anesthesiology, General Hospital,
Respiratory, Infection Control and
Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indications for Use

510(k) Number (if known)
K130800

Device Name
Powder Free Vinyl Patient Examination Gloves, Clear (Non-colored)

Indications for Use (Describe)
Powder Free Vinyl Patient Examination Gloves, Clear (Non-colored) is a non-sterile disposable device intended for medical purposes that is worn on the examiner's hand or finger to prevent contamination between patient and examiner.

Type of Use (Select one or both, as applicable)

☐ Prescription Use (Part 21 CFR 801 Subpart D)

☒ Over-The-Counter Use (21 CFR 801 Subpart C)

PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON A SEPARATE PAGE IF NEEDED.

FOR FDA USE ONLY

Concurrence of Center for Devices and Radiological Health (CDRH) (Signature)

Elizabeth F. Claverie-S
2014.02.15 00:27:45-05:00'

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